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TITLE

METHOD FOR COPPER PLATING

ABSTRACT :

PURPOSE: To obtain a copper plating layer of good adhesive strength by electrolytically pickling a steel material in an aq. sulfunc acid which contg. thiourea or its deriv. without using cyanide harmful for sanitation then electroplating the same in a copper sulfate bath.

CONSTITUTION: About 0.005~0.5g/lthiourea or deriv. of thiourea such as anyl thiourea is added into an aq. sulfuric acid contg. about 20~100g/l sulfuric acid, and with a steel material as cathode, an electrolytic pickling treatment is accomplished at about 10~40°C, about 10~500A/dm² current density and for about 0.5~10sec electricity conduction time. Immediately after this without washing by water, the steel material is electroplated with a copper sulfate plating bath similar to that by known methods. The thiourea or its deriv. electrostatically charges to ⊕ in the pickling bath, sticks and coats the surface of the steel material electrically in the electrolytic pickling stage, prevents replacement plating reaction in the electroplating stage and improves the adhesive strength of the copper plating layer considerably.

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69銅めつき方法

②特

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22出

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明細響

1.発明の名称

銷めつき方法

2.特許請求の範囲

(1)鋼素地上へ銅めつき屬を形成するに当り、チオ尿素又はその誘導体を含有する硫酸水溶液中で、被めつき鋼材を陰極として電解酸洗を行なつた後、硫酸銅浴中で電気めつきを行なうととを特徴とする銅めつき方法。

3.発明の詳細な説明

本発明は網めつき方法に関し、殊に硫酸網浴を 用いて頻繁地の表面に密育性の良好を網めつき層 を形成する方法の改良に関するものである。

従来、硫酸網浴を用いて密着性の良好な網のつき層を形成する為には、硫酸網電気めつきに先立つてシアン化網浴によるストライクめつきを行なうととが不可欠とされていた。しかしながらこの工程は、シアン化物を含有する排液公害を生じるという重大な問題があり、かかる問題を回避する為にはシアン化合物を用いない網めつき法の確立

が必要で、多くの研究が行なわれている。

これら改良研究の殆んどは、就酸網浴中での電気めつき工程で超こる網と網との関携めつき反応を抑制し、網めつき層の密管性低下を助止しようとするものである。例えば特公昭 87-5708号には、酸鉄抑制剤を含む酸洗浴で被めつき材を予備処理した後硫酸鍋浴中で電気めつきを行なうことにより、密管性の良い鍋めつき層が得られると報告されている。また特公昭 5 4 - 4 8 2 9 号は、硫酸網浴に置換抑制剤としてアリルーチオ原素を添加し、 般換めつき反応を抑制しつつ電気めつきを行なうことによって、網めつき層の密管性を高めようとするものである。

本発明者等もかれてよりシアン化合物を使用しない網めつき法の開発を期して種々研究を行えつているが、その研究の過程で前記公告公報配収の方法を追試した。その結果、鍋めつき 層の密度性に関しては、記載された程の効果が得られなかつた。 そとでその理由を退求したととろ、以下に示す様な事実が確認された。即ち上記公報記載の方

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特開昭57-116799(2)

又はその誘導体を添加しておき、彼めつき材を触 **極として電解酸洗を行なりが、チオ原素やその誘** 導体は硫酸水溶液中で⊕に帯電しているから、質 解酸洗工程では浄化された被めつき材の表面に置 気的に引き寄せられ、彼めつき材の表面全域を完 全に被覆する。従つてこの予備処理の後硫酸銅浴 に違いて電気めつきを行なうと、全面質気めつき からなる密管性の優れた網めつき間を得ることが できる。この場合、予備処理後は水洗することな くそのまま憶気めつきを行なりのがよい。尚削記 公告公翻記載の方法においても、酸値抑制剤又は 置換抑制剤としてチオ尿器やその誘導体を使用で きる盲記載されている。しかしこれらの方法では、 上記の楽剤を酸洗浴又は監換網めつき浴等に添加 して微換めつき反応を抑制しようとするものであ り、これらの製剤の彼めつき材表面への付譲は物 理的な付借のみであるから、チオ尿繁等を多量流 加して被役刑決度を高める必要があると共に、仮 に 沸度を十分に高めた場合でも、ミクロ的 見地か らみると完全な被覆効果は得られない。また依解

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法は、被めつき材の表面全域を酸蝕抑制和又は懺換抑制剤で被覆することによつて懺換めつき反応を抑制しようとするものであるが、ミクロ的見地からすると被殺効果が不十分であつて質換めつき反応が起こって当該部分のでは、部分的には懺換めつき反応が起こって当該部分のでは、とかできる。従って電気めつき不能を完全に被破する。とかでき、且つ電気めつきを阻害しない様を下する。ないでき、且つ電気めつきを阻害しない様をあめることができると考えた。

本発明はかかる知見を基に、予備処理法の改善 によつて電気網めつき層の密着性を高めるべく鋭 意研究の結果完成されたものであつて、その糖放 は、網累地上へ鍋めつき層を形成するに当り、チ オ尿緊又はその誘導体を含有する硫酸水溶液中で、 被めつき網材を陰極として電解酸洗を行なつた後、 硫酸網浴中で電気めつきを行なりととろに襲音が 存在する。

本発明では、酸洗用の硫酸水溶液中にチオ尿繁

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めつき浴中にチオ尿素等を添加し、電解めつき工 程で同時にチオ原業等による表面被變効果を発揮 させて厳拠めつきを抑制する方法も考えられる。 しかしとの方法では、電気めつきの初期段階で慌 換めつき反応が併発し、密育性はやはり不十分に なる。これらに対し本発明の方法では、網イオン の全く存在しない酸洗処理の段階で、被ぬつき材 の表面全域にサオ尿素等を質気的に付着させて被 覆する方法であるから、比較的低濃度のチオ原絮 等であつても十分な効果を得ることができ、被覆 効果も殆んど完全無欠である。更に上記公告公報 記載技術の改善法として、アセチルチオ尿薬を含 む酸洗浴に被めつき材を浸漬して酸洗した後、ア セチルチオ尿素を含む硫酸銅浴で置換めつきを行 ない、水洗後硫酸銅浴中で電気めつきを行なり方 法が報告されている。しかしこの方法は予備処理 が煩雑であると共に、微換めつき工程での浴組成 が催かに違うだけで密滑性が振端に低下する傾向 があり、しかもアセチルチオ尿素の付着は前配と 同様所栓物理付着のみであるから、置換めつき反

応抑制効果は本発明法に比べてはるかに劣る。

本発明で便削するチオ原架又はその誘導体は、 前述の如く酸洗浴中で⊕化帮制し、電解酸洗工程 で歌気的に被めつき材影面に付着して漫画被舞効 泉を発排するもので、チオ尿絮誘導体としては、 アリルチオ尿器、アセチルチオ尿器、N-アリル - N- アルキルチオ尿器、N-アルアルキル- N - Tルキルチオ尿素、N-シクロアルキル- N-アルキルチオ尿器、N-アルキル-N-アルキル チオ尿絮、B-アルキルチオ尿器及びそのN-ア ルキル艦換体、N-フエニル-N-オキシエチル チオ泉系、N-シクロヘキシルーN-オキシエチ ルチオ尿素、N-アルキル-N-オキシエチルチ オ泉紫、N-フエニル-N-オキシプロピルチオ 尿素、N-フエニル-N-β-(オキシエトキシ) - エチルチオ尿紫、N-ペンジル- N'- オキシエ チルチオ尿素、N-4-オキシフエニル-N-オ キシエチルチオ尿紫等が例示される。上記誘導体 中のアルキル残酷としては、炭素数が6以下の低 級アルキル基が好ましい。これらの中でも最も優

れた効果を発揮するのはナオ尿素、アリルチオ尿 繁及びアセチルーチオ尿素である。 尚後記実験例 でも明らかにする如く、チオ尿素等と同機酸洗浴 中で①に帯電するアミン系化合物を使用しても電 解めつき腐の密着性は全く改善されず、本発明者 等が実験した限り前述の様な優れた密音性向上効 果を発揮するのはチオ尿素及びその誘導体のみで あつた。

電解酸洗浴として使用する硫酸水溶液の濃度は特に限定されないが、最も好ましいのはその1 & 中に硫酸20~100%を含有するものであり、チオ尿薬又はその誘導体はこの水溶液中に0.005~0.5%/8、より好ましくは0.01~0.2%/8 添加される。電解酸洗処理は、上記酸洗浴(好ましくは10~400程度)中で、绷素地を除版として通難することによつて行なわれるが、このともの最も好ましい電流密度は10~500A/dm²、通難時間は0.5~10 勘程度である。尚鋼架地は予め脱脂しておくのがよい。この様にして観解したおくのがよい。この様にして観解したそうと、鋼器地の表面金数にチオ保熱又は

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機硫酸を609/6含有する硫酸水溶液に0.19/6のチオ尿素を加えて電解酸洗液を調製し、この浴に脱脂処理した鋼線材を浸漬走行させながら、電流密度800Å/dm²で1秒間の電解酸洗を行なり。この予備処理線材を水洗することなく硫酸銅めつき浴(Cu8O4・5H2O:8009/6、漁田28O4:409/6)に導き、電流密度200Å/dm²、線材走行速度:250m/分で2秒間電気めつきを行ない、得られた鯛めつき鋼線材の密體性及びめつき液中への鉄の溶出量を調べた。尚密管性は、得られた鯛めつき鋼線10本を共巻き試験に付し、鋼めつき層に全く剝離が認められないものの数で判断した。

また電解酸洗液にチオ尿業を添加しなかつた他 は上記と同様にして電解酸洗及び電気めつきを行 ない、密着性及び鉄溶出塩(被めつき材の単位表 面積当りの溶出量)を購べた。

結果を第1表に示す。



特別昭57-116799(3)

その誘導体が付着するから、その後硫酸的めつき 浴に導いて電気めつきを行なえばよい。電気めつ き処理は公知の方法と全く同様にして行なえばよ く、例えば水1 ℓ 当り硫酸銅170~850 ₽を 含有する硫酸網めつき浴を用い、温度10~80 で、電流密度5~800 A/d m² で通電すること によつて行なわれる。この場合硫酸網めつき浴に も少量のチオ尿紫又はその誘導体を添加すること ができる。

本発明は機略以上の際に構成されており、チオ 尿素又はその誘導体を含む酸洗浴を用いた電解酸 洗法を採用することによつて、電気めつき工程で の置換めつき反応を防止し網めつき溜の密着性を 大幅に高め得ることになつた。しかも本発明では、 鋼繁地の酸洗工程で簡時に罹鈍めつき防止用予備 処塊を行なりものであるから、予備処理工数が増 加したり処理時間が延長する等の問題もなく極め て実用に即した方法と言える。

次に契験例を示す。

突験例I

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第	1		
チオ尿薬激派の有無	铅着性	鉄溶出量 (g/m²)	備考
有	10/10	0, 5	與施例
無	6/10	1. 7	比較例

第1 表からも明らかな様に、本発明法(契額例)ではめつき液への鉄器出量が極めて少なく密治不良のものは皆無であるが、チオ尿素の張加を省略した比較例ではめつき液への鉄路出量が多く驚懊めつき反応が起こつており、4 割もの密機不良品が発生している。

突験例2

結果を第2表に一括して示す。

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第 2 数

愷	換めつき防止剤	·	密療性	鉄容出數	備考
添加場所	箱 類	添加進	120 MM 124	(9/m²)	雅考
電 解 酸 洗 液	チオ尿素	0.1 8 / 6	10/10	0. 5	実
	アリルチオ尿素	0.1 9 / e	10/10	0.8	165
	アセチルチオ尿素	0.1 8 / 8	10/10	0. 4	例
	アミンA	100/8	0/10	1.1	比
	т ぇ ン B	100/8	0/10	1. 1	較
	アミンロ	100/6	0/10	1. 0	例
電気めつき液	チオ原素	0.1 9 / 2	7/10	0. 7	
	アリルチオ尿紫	0.14/6	7/10	0.6	
	フセチルチオ尿素	0.18/6	7/10	1.0	叁
	アミンム	100/8	0/10	1, 4	考
	アミンB	100/6	0/10	1. 8	<i>19</i> 1
	アミンロ	100/6	0/10	1. 8	
	無添	ŻNI	6/10	1. 7	Ī

アミンA:イピツトル B-600(住友化学社製商品名) アミンB:イピツトル B00-LA(住友化学社製商品名) アミンC:イピツトル 5 B 0 (住友化学社製商品名)

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突躁例4

即ち相当量のチオ界緊を酸洗液に添加した場合 であつても、電解酸洗法を採用しない限り本発明 の目的性違成できない。

4.図面の簡単な説明

第1~8 圏は、チオ尿絮、アリルーチオ尿絮及 びアセチルーチオ尿絮の添加量と皆着性の関係を 示すグラフである。

> 出願人 株式会社神戸製鋼所 代理人 弁理士 橋 木 久 一般出

第2表からも明らかな様に、アミン化合物は電解酸洗液中で砂に帯電していると考えられるが、その密菌性破醫効果は全く認められず、無添加のものよりもはるかに思い。またチオ尿素等は電解酸洗液に添加したときに限つて卓越した密層性改善効果を発揮しており、電気めつき液のみに添加しても十分な効果は得られない。

実験例8

置機めつき筋止剤としてチオ尿素、アリルチオ 尿素及びアセチルチオ尿素を選択し、実験例1の 方法に挙じて失々の添加量と格管性の関係を調べ た。

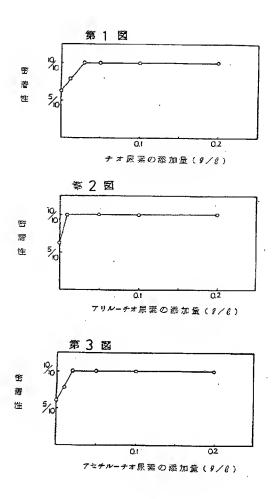
結果を第1國(チオ尿薬)、第2図(アリルチオ尿薬)及び第8図(アセチルチオ尿薬)に夫々示す。

第1~8 図からも明らかな際化、十分な密衛性 を確保する為には、チオ尿繁で 0.0 8 9 / 8 以上、 アリルチオ尿繁で 0.0 1 8 / 8 以上、アセチルチ オ尿素で 0.0 2 9 / 8 以上を失々復解酸洗液中化 添加することが領まれる。

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特開 昭57-116799 (5)



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- 15 Request for Examination: not filed (5 pages in total)
 - (54) Title of the Invention: COPPER PLATING METHOD
 - (21) Application Number: S56-2264
- 20 (22) Application Date: January 9, 1981
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SPECIFICATION

1. TITLE OF THE INVENTION

35 COPPER PLATING METHOD

2. CLAIMS

A copper plating method for forming a copper plating layer on a steel base material, comprising

a first step of electrolytic pickling a steel material to be plated in a sulfuric acid aqueous solution containing thiourea or a thiourea derivative, the steel material being the cathode, and

a second step of electroplating the steel material in a copper sulfate bath.

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3. DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to a copper plating method, especially an improved version of a method for forming a copper plating layer with excellent adhesive property on the surface of a steel base material by using a copper sulfate bath.

Conventionally, in order to form a copper plating layer with excellent adhesive property by using a copper sulfate bath, it has been indispensable to perform strike-plating in a copper cyanide bath prior to the copper sulfate electroplating. This process, however, problematically causes pollution due to the waste liquid containing cyanides. It has therefore been desired to establish a copper plating method in which a cyanide is not used, and many researches have been conducted to achieve this goal.

Most of those researches aim to suppress displacement plating reaction between the copper and the steel that occurs in the electroplating process in the copper sulfate bath so as to prevent deterioration of the adhesive property of the copper plating layer. For example, JP-B S37-5708 has reported that a copper plating layer with excellent adhesive property can be obtained by preliminary processing a material to be plated in a pickling bath containing an pickling inhibitor and then electroplating the material in a copper sulfate bath. Moreover, JP-B 54-4329 discloses an invention aiming to improve the adhesive property of a copper plating layer by adding

allylthiourea as displacement inhibitor in a copper sulfate bath so as to suppress displacement plating reaction and then performing electroplating.

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The present inventors have conducted various researches to develop a copper plating method in which a cyanide is not used. During the researches, the present inventors conducted supplemental examinations of the methods disclosed in the foregoing patent publications. As a result, it was found that the effects on the adhesive property of the copper plating layer were not as much as those disclosed in the documents. present inventors pursued the reasons and found the following. Namely, according to the methods of the above patent publications, the displacement plating reaction is inhibited by coating the whole surface of the material to be plated with a pickling inhibitor or a displacement inhibitor. From a microscopic point of view, however, the effect of the coating is insufficient for fully inhibiting the displacement plating reaction. Therefore, the displacement plating reaction occurs in some parts so that the adhesive property of the parts deteriorates. Based on the findings, the present inventors assumed that it was possible to improve the adhesive property of the electroplating layer by establishing a preliminary processing method in which the whole surface of a material to be plated can be fully coated with the foregoing agents, and accordingly the electroplating may not be inhibited.

The present invention has been devised as a result of the intensive investigations to increase the adhesive property of the electroplating layer by improving the preliminary processing method based on the foregoing findings. The gist of the present invention is to firstly carry out electrolytic pickling of steel material to be plated in a sulfuric acid aqueous solution containing thiourea or a thiourea derivative, the steel material being the cathode, and then electroplating the steel material in a copper sulfate bath, in the process of forming a copper plating layer on the steel base material.

According to the present invention, thiourea or a thiourea derivative is added to an aqueous sulfuric acid solution for pickling, and electrolytic pickling of the material to be plated is performed with the material as the cathode. In this process, as the thiourea or the thiourea derivative is positively charged in the aqueous sulfuric acid thiourea or the thiourea derivative is solution, the electrically drawn to the surface of the material to be plated which has been washed in the electrolytic pickling process so as to fully coat the whole surface of the material to be plated. 10 Accordingly, when the resulting material to be plated after the preliminary processing is subjected to electroplating in a copper sulfate bath, the whole surface of the material can be electroplated with a copper layer with excellent adhesive property. In this process, it is preferable to perform the 15 electroplating without water-washing after the preliminary Meanwhile, use of thiourea or a thiourea processing. derivative as a pickling inhibitor or a displacement inhibitor is described in the methods of the aforementioned patent However, according to those methods, the publications. 20 inhibitor is added to the pickling bath or the displacement copper plating bath so as to inhibit displacement plating reaction. Thus, as the adhesion of the agent to the material to be plated is based only on physical adhesion, it is necessary 25 to increase the concentration of the coating agent by adding a large amount of thiourea or similar agents. Furthermore, even if the concentration is sufficiently increased, a perfect coating effect cannot be obtained from a microscopic point of view. It may be possible to exemplify another method for inhibiting displacement plating, in which thiourea or a similar 30 agent is added to an electroplating bath so that the surface coating effect of the thiourea or the similar agent is achieved In this method, however, a in the electroplating bath. displacement plating reaction concomitantly occurs at an early stage of the electroplating, resulting in insufficient adhesive 35

property. On the other hand, according to the method of the present invention, the whole surface of the material to be plated is electrically coated with thiourea or a similar agent at a stage of the pickling process where no copper ions exist. For this reason, it is possible to achieve sufficient effects even with the thiourea or the similar agent having a relatively low concentration, and furthermore, the resulting coating effect is almost perfect. Furthermore, in order to improve the techniques described in the aforementioned patent publications, it has been reported a method in which: a material to be plated 10 is immersed in a pickling bath containing acetylthiourea for electrolytic pickling; the material to be plated is then subjected to displacement plating in a copper sulfate bath containing acetylthiourea; water-washing is performed; and the resulting material is electroplated in a copper sulfate bath. 15 This method, however, has problems; namely, the preliminary processing is complicated, and even a slight change in the composition of the bath tends to extremely decrease the adhesive property of the resulting product. Additionally, since the acetylthiourea is only physically adhered similarly as the 20 above case, the effects of inhibiting displacement plating reaction is far inferior to the method according to the present

The thiourea or a thiourea derivative used in the present 25 invention is positively charged in the pickling bath as mentioned earlier and is electrically adhered to the surface of the material to be plated to exert the surface coating effect. Examples of the thiourea derivative include allylthiourea, N-ally-N'-alkylthiourea, acetylthiourea, N-aralkyl-N'-alkylthiourea, N-cycloalkyl-N'-alkylthiourea, 30 N-alkyl N-alkyl-N'-alkylthiourea, 8-alkylthiourea and substituted analogs thereof, N-phenyl-N'-oxyethylthiourea, N-cyclohexyl-N'-oxyethylthiourea, N-phenyl-N'-oxypropylthiourea, $N-phenyl-N'-\beta-(oxyethoxy)-ethylthiourea,$ 35

invention.

N-benzyl-N-oxyethylthiourea, and N-4-oxyphenyl-N'-oxyethylthiourea. Examples of the alkyl residue in the derivative preferably include lower alkyl groups having 6 or less carbons. The examples showing the most excellent effects among the above examples are thiourea, allylthiourea, and acetyl-thiourea. As will be clearly shown by Examples mentioned below, in the case where an amine compound, which is positively charged in the pickling bath similarly as thiourea, is used, the adhesive property of the resulting electroplating layer is not at all improved. Based on the examination results conducted by the present inventors or others, it is only thiourea or a thiourea derivative that can exert excellent adhesive property-improving effects mentioned earlier.

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The concentration of the aqueous sulfuric acid solution to be used as an electrolytic pickling bath is not particularly limited. The most preferable example is an aqueous sulfuric acid solution containing 20 to 100 g of sulfuric acid in one liter of the solution, in which thiourea or a thiourea derivative is contained in an amount of 0.005 to 0.5 g/l, or preferably 0.01 to 0.2 g/l. The electrolytic pickling is performed by supplying electricity to the steel base material as the cathode in the aforementioned pickling bath (preferably at a temperature of 10 to 40°C). In this process, the electric current density is preferably 10 to 500 A/dm² and the electricity supplying time is preferably about 0.5 to 10 seconds. The steel base material is preferably degreased in advance. When the electrolytic pickling is performed in the aforementioned manner, the thiourea or the thiourea derivative is adhered to the whole surface of the steel base material. Thereafter, the material may be introduced to a copper sulfate plating bath for electroplating. The electroplating may be performed in the same manner as conventional methods, for example, by using a copper sulfate plating bath containing 170 to 850 g of copper sulfate per one liter of water and supplying electricity at a

temperature of 10 to 80° C and with the current density of 5 to 300 A/dm^2 . It is possible to add a small amount of the thiourea or the thiourea derivative in the copper sulfate plating bath.

The present invention with the above-outlined structure can greatly improve the adhesive property of the copper plating layer by employing the electrolytic pickling method in which the pickling bath contains thiourea or a thiourea derivative and thus displacement plating reaction can be prevented from occurring in the electroplating process. Moreover, according to the present invention, the preliminary processing for preventing displacement plating is simultaneously performed during pickling of the copper base material, and thus problems such as increase of the number of preliminary processing and a longer processing time do not occur. Therefore, the present invention is extremely useful and practical.

Examples of the present invention are described below.

Example 1

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An electrolytic pickling solution was prepared by adding 0.1 g/l of thiourea in an aqueous sulfuric acid solution containing 60 g/l of concentrated sulfuric acid. By immersing and running a degreased steel wire in the thus obtained electrolytic pickling bath, the wire was electrolytic pickled with the current density of 800 A/dm2 for one second. preliminary processed steel wire was introduced into a copper sulfate plating bath (CuSO₄·5H₂O: 300 g/l, concentrated H₂SO₄: 40 g/l) without water washing, and subjected to electroplating for two seconds (current density: 200 A/dm2, running speed of steel wires: 250 m/min.). The resulting copper-plated steel wire was examined for the adhesive property and the amount of iron dissolved in the plating bath. By subjecting ten wires of the thus obtained copper-plated steel wires to co-winding test, the adhesive property of the plated layer was determined based on the number of the wires with no peeling in the copper plated layer.

Additionally, electrolytic pickling and electroplating were performed in the same manner as described above, except that thiourea was not added in the electrolytic pickling solution. The adhesive property and the amount of dissolved iron (amount of dissolved iron per unit surface area of the plated material) were measured. Table 1 shows the results.

Table 1

Thiourea	Adhesive property	Dissolved iron (g/m²)	Remarks
Added	10/10	0.5	Example
Not added	6/10	1.7	Comparative Example

As is evident from Table 1, in the case of the method according to the present invention (Example), the amount of iron dissolved in the plating bath was extremely small and there was no case in which the adhesive property of the plated layer was defective. On the other hand, in the case of Comparative Example in which thiourea was not added, the amount of iron dissolved in the plating bath was large; displacement plating reaction occurred; and as much as 40 percent of the plated layer had a defect in terms of the adhesive property.

20 Example 2

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Electrolytic pickling and electroplating were performed in the same manner as in Example 1 while using different kinds of displacement plating inhibitors, and the adhesive property of the copper plating and the amount of iron dissolved in the electroplating bath were examined. For reference, the effect obtained when the displacement plating inhibitor was added in the electroplating bath was examined.

Table 2 shows all the results.

Table 2

Displacem Addition target	nent plating inhib	Addition amount	Adhesive property	Dissolved Iron (g/m²)	Remarks	
Electrolytic	Thiourea	0.1 g/l	10/10	0.5	Example	
pickling	Allylthiourea	0.1 g/l	10/10	0.8		
solution	Acetylthiourea	0.1 g/l	10/10	0.4		
	Amine A	1 cc/l	0/10	1.1	Componetizza	
	Amine B	1 cc/l	0/10	1.1	Comparative Examples	
	Amine C	1 cc/1	0/10	1.0		
Electroplating	Thiourea	0.1 g/l	7/10	0.7		
solution	Allylthiourea	0.1 g/l	7/10	0.6		
	Acetylthiourea	0.1 g/l	7/10	1.0	Reference	
	Amine A	1 cc/l	0/10	1.4	Examples	
	Amine B	1 cc/l	0/10	1.8		
	Amine C	1 cc/l	0/10	1.8		
No addition		6/10	1.7			

Amine A: Ibit No. 8-600 (Product name, Sumitomo Chemical Co., Ltd.)

Amine B: Ibit No. 600-LA (Product name, Sumitomo Chemical Co., Ltd.)

5 Amine C: Ibit No5808-600 (Product name, Sumitomo Chemical Co., Ltd.)

As is evident from Table 2, although assumably the amine compounds were positively charged in the electrolytic pickling solution, there was no sign of the adhesive property-improving effects and further the adhesive properties were far inferior to that in the case where none of the agents was added. Moreover, thiourea or a similar compound showed excellent adhesive property-improving effects only when it was added in the electrolytic pickling agent, and thus a sufficient effect could not be obtained when it is added only in the electroplating solution.

Example 3

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Thiourea, allylthiourea, and acetylthiourea were

selected as the displacement plating inhibitor, and the relations between each of the addition amounts and the adhesive property were examined according to the method described in Example 1.

Fig. 1 (for thiourea), Fig. 2 (for allylthiourea) and Fig. 3 (for acetylthiourea) show the results.

As is evident from Figs. 1 to 3, in order to achieve sufficient adhesive property, it is necessary to add thiourea in an amount of not less than 0.08 g/l, or allylthiourea in an amount of not less than 0.01 g/l, or acetylthiourea in an amount of not less than 0.02 g/l in the electrolytic pickling solution.

Example 4

Copper plating was performed in the same manner as in Example 1, except that 0.1 g/l of thiourea was added in the pickling solution and a simple immersion-pickling method was employed without power distribution. The adhesive property of the resulting copper-plated steel wire was 8/10, which indicates that the resulting product has a 20% chance of having the adhesive defect.

Namely, even if a considerable amount of thiourea is added in the pickling solution, the goal of the present invention cannot be achieved unless the electrolytic pickling method is employed.

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4. BRIEF DESCRIPTION OF THE DRAWINGS

Figs. 1 to 3 each illustrates a graph showing relations between the adhesive property and the addition amount of thiourea, allylthiourea or acetylthiourea.

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Fig. 1

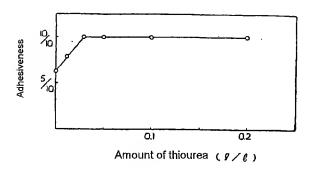
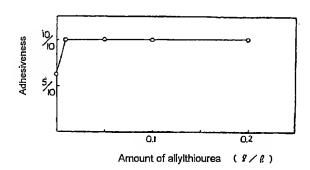
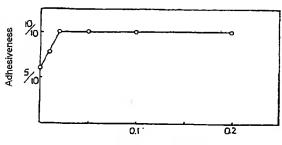


Fig. 2



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Fig. 3



Amount of acetylthiourea (9/0)